

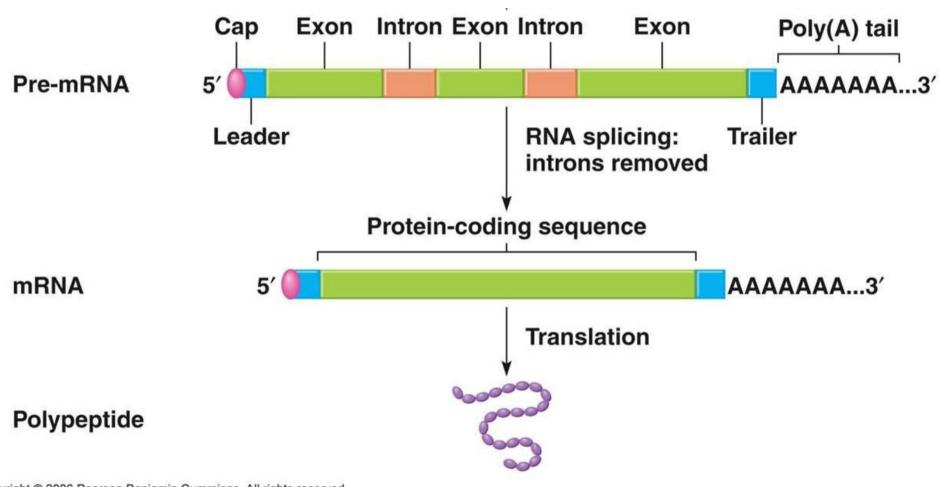
# **RNA Structure Genetic Code Translation**

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# Type of RNA

- 1. Ribosomal RNA (rRNA)
  - 16S (small ribosomal subunit)
  - 23S (large ribosomal subunit)
  - 5S (large ribosomal subunit)
- 2. Transfer RNA (tRNA)
- 3. Messenger RNA (mRNA)

# hn-RNA (Pre-mRNA) & m-RNA



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# m-RNA

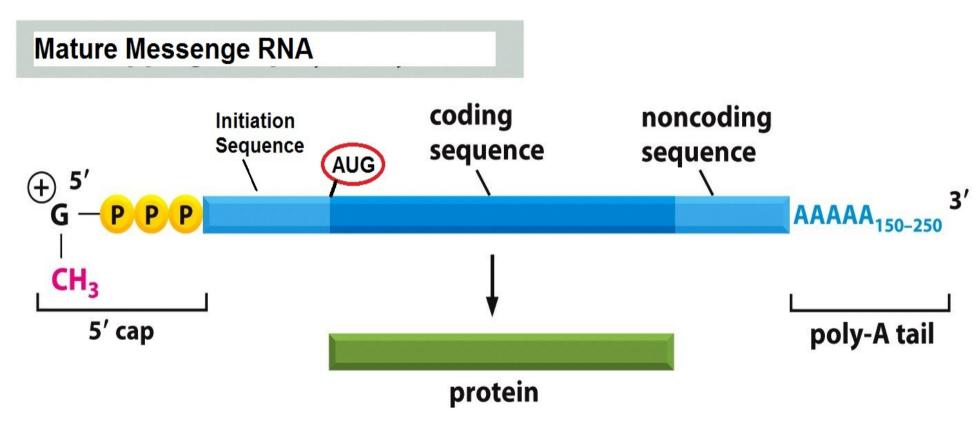
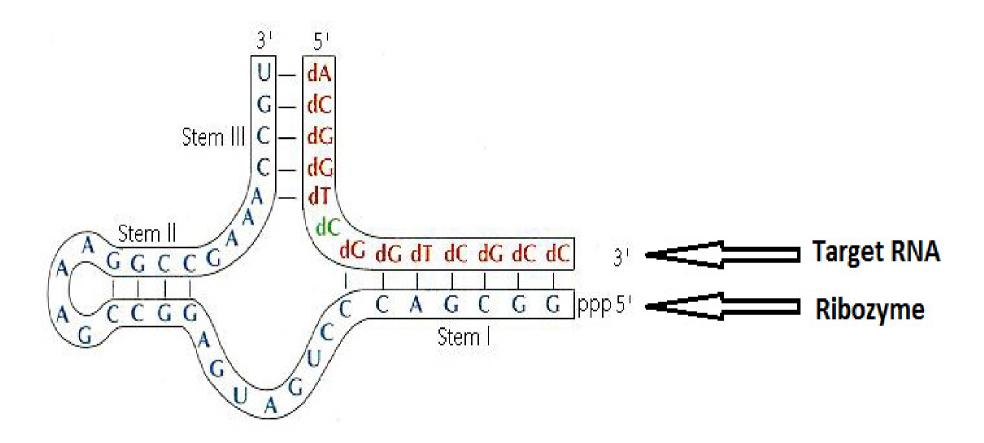
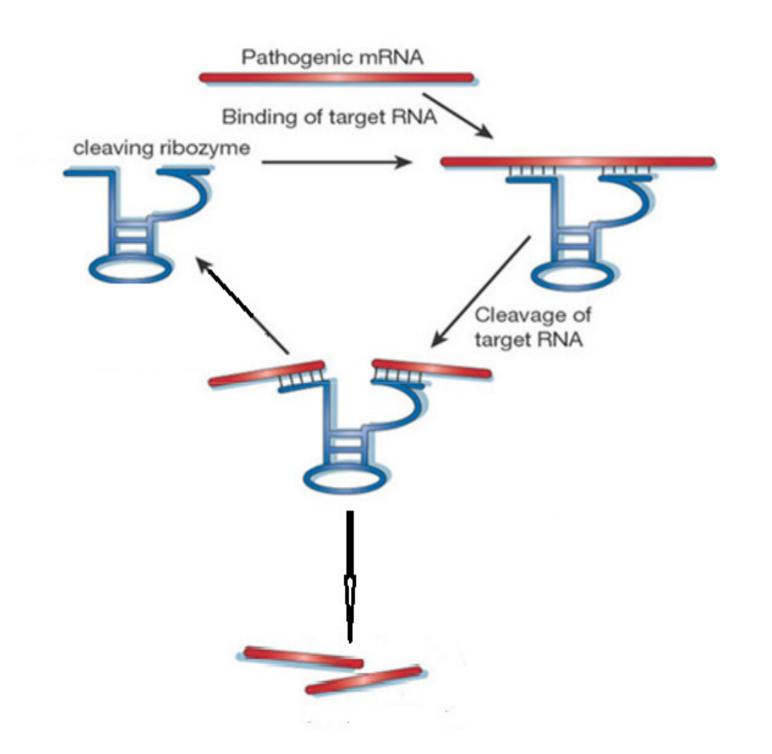


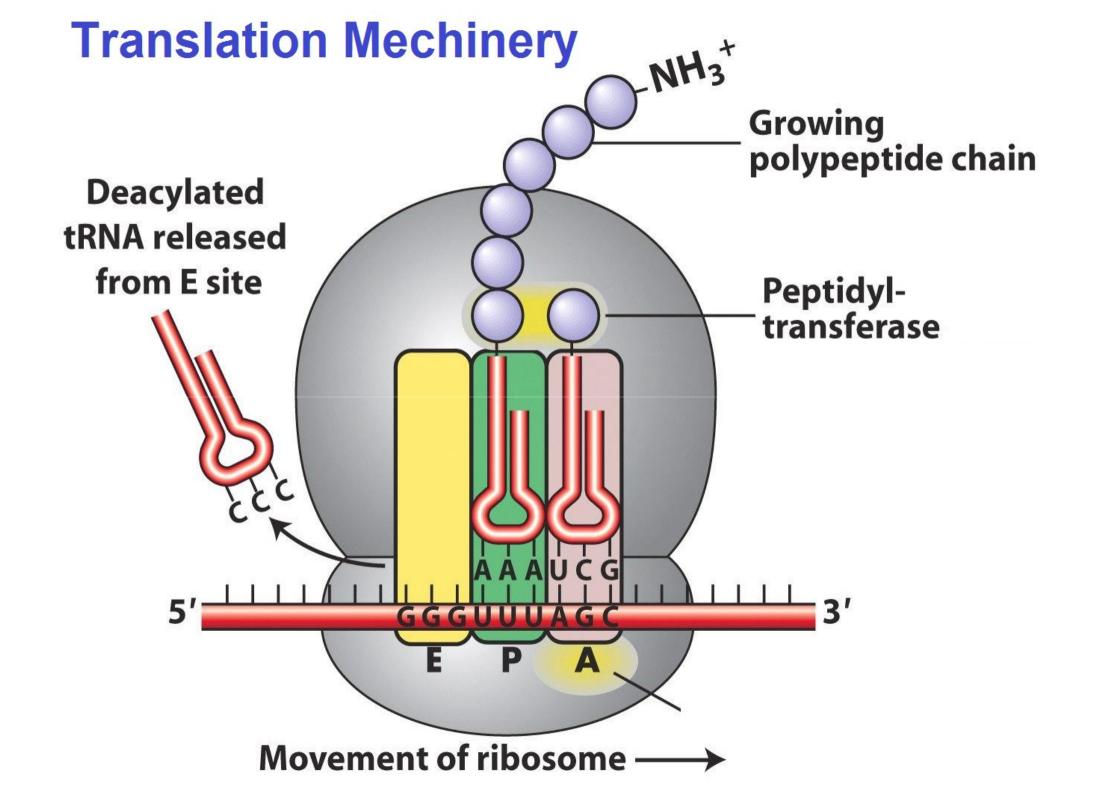
Figure 7-16a Essential Cell Biology 3/e (© Garland Science 2010)

# r-RNA (Ribosomal RNA)

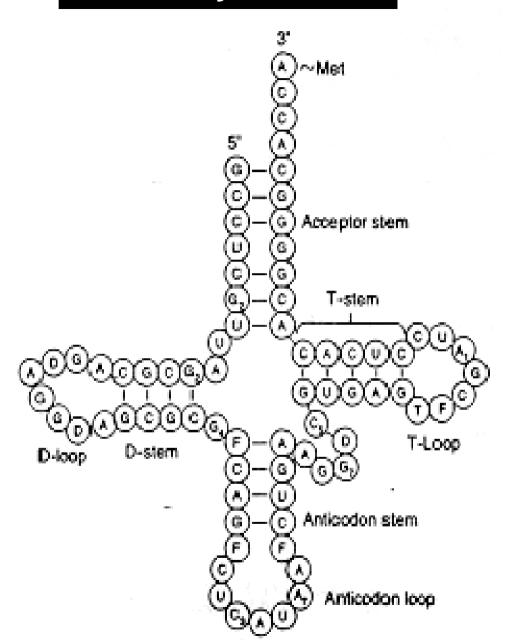
- Association with several proteins
- Type :
  - > In Prokaryots = 23S, 16S, and 5S
  - > In Eukaryots = 28S, 18S, 5.8S, and 5S
- "S" = Svedberg unit
   Related to the molecular weight and shape.
- Function :
  - Sites for protein synthesis.
  - Catalysts in protein synthesis.
    - E.g. "Ribozyme".

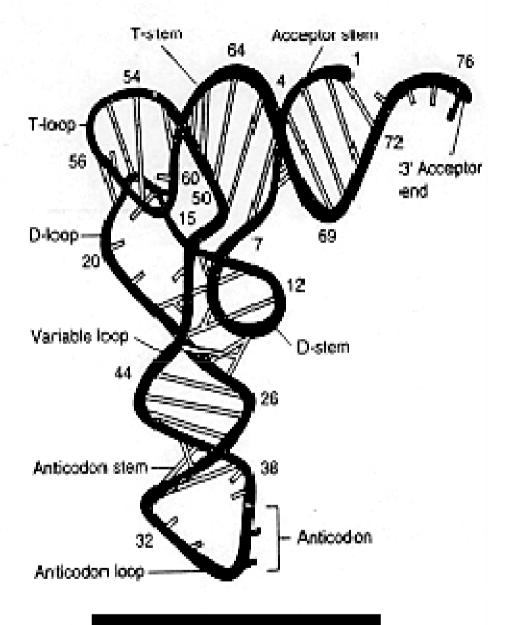






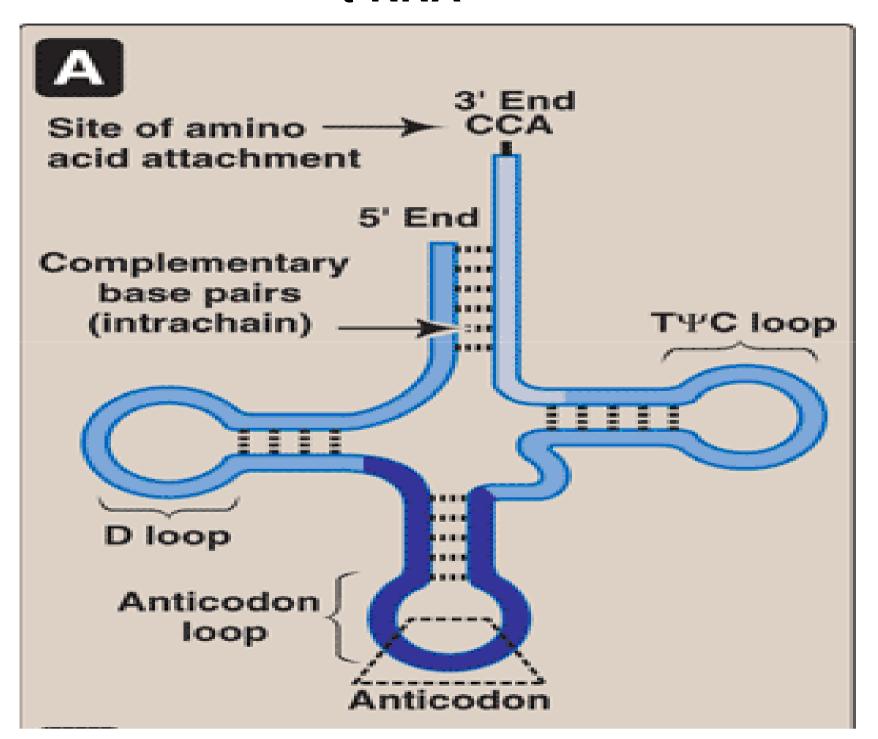
### **Secondary structure**





**Tertiary structure** 

### t-RNA

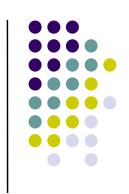


# t-RNA (transfer RNA)

- Smallest (73 93 ns)
- Easily soluble = s-RNA
- Specific tRNA for All 20 amino acids.
- Clover leaf like structure.
- Unusual bases (for example, dihydrouracil,)
- Intrachain base-pairing = Looks secondary & tertiary structure.
- Serves as an "adaptor" molecule

- Dihydrouracil arm = Recognition Enzyme to add amino acid
- Pseudouridine arm = Binding t-RNA to ribosome
- Anticodon arm = Recognize triplet codon on m-RNA
- Accepter arm = carries amino acid

# **Genetic Code**



- Nucleotides read in triplet "codons"
  - 5'  $\rightarrow$  3'
- Each codon translates to an amino acid
- 64 possible codons
  - 3 positions and 4 possiblities (AGCU) makes 4<sup>3</sup>
     or 64 possibilities
  - Degeneracy or redundancy of code
    - Only 20 amino acids
    - Implications for mutations

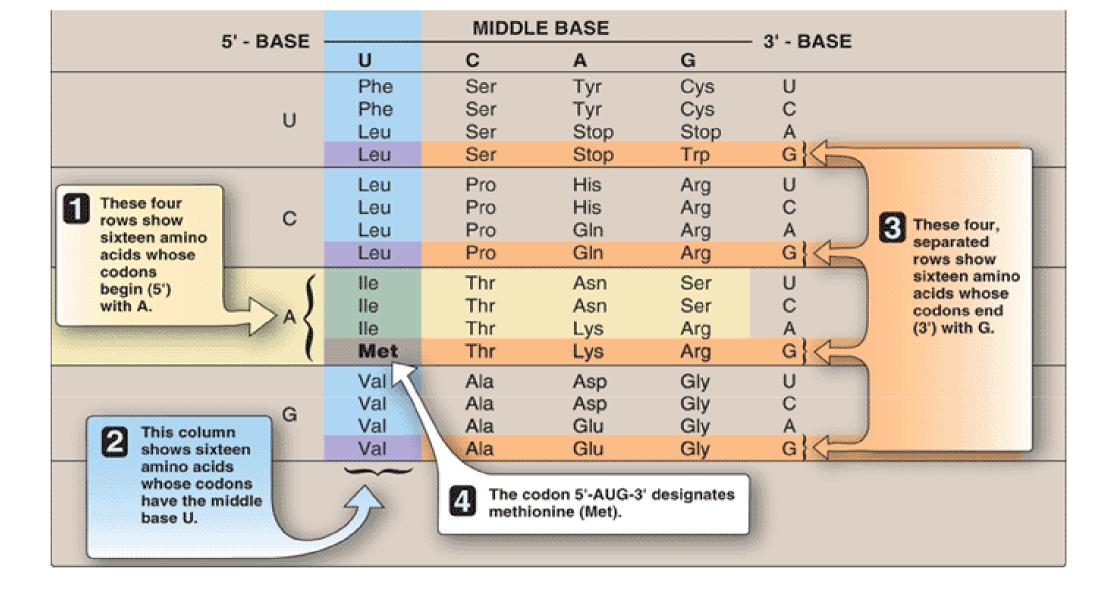
# **Genetic Code**

first position



### second position

	U	С	A	G
U	UUU Phe (F) UUC Phe (F) UUA Leu (L) UUG Leu (L)	UCC Ser (S) UCA Ser (S)	UAU Tyr (Y) UAC Tyr (Y) UAA STOP UAG STOP	UGC Cys (C)
С	CUC Leu (L) CUA Leu (L)	CCC Pro (P) CCA Pro (P)	CAU His (H) CAC His (H) CAA Gln (Q) CAG Gln (Q)	CGC Arg (R) CGA Arg (R)
A	AUU Ile (I) AUC Ile (I) AUA Ile (I) AUG Met (M)		AAC Asn (N)	AGC Ser (S)
G	GUC Val (V)	• •		GGC Gly (G)



# Use of the genetic code table to translate the codon AUG

### **Genetic Code**

### 1. Triplet Codons:

Each codon is a consecutive sequence of three base

### 2. Non-overlapping:

Codes are always read one after another.

### 3. Non-punctuated:

Codes are always continues.

# 4. Specificity (unambiguous):

One codon always codes for specific amino acid

### 5. Universal:

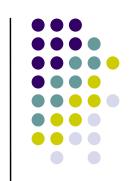
Codons represent same amino acid in all species.

Except in Mitochondria



### 6. Degenerate:

Each codon corresponds to a single A.A. But A.A. may have more than one triplet codon.



### 7. Initiator codon:

AUG is start codon

### 8. Termination codons

Nuclear DNA = UAA, UGA, UAG, Mitochondrial DNA = AGG, AGA,

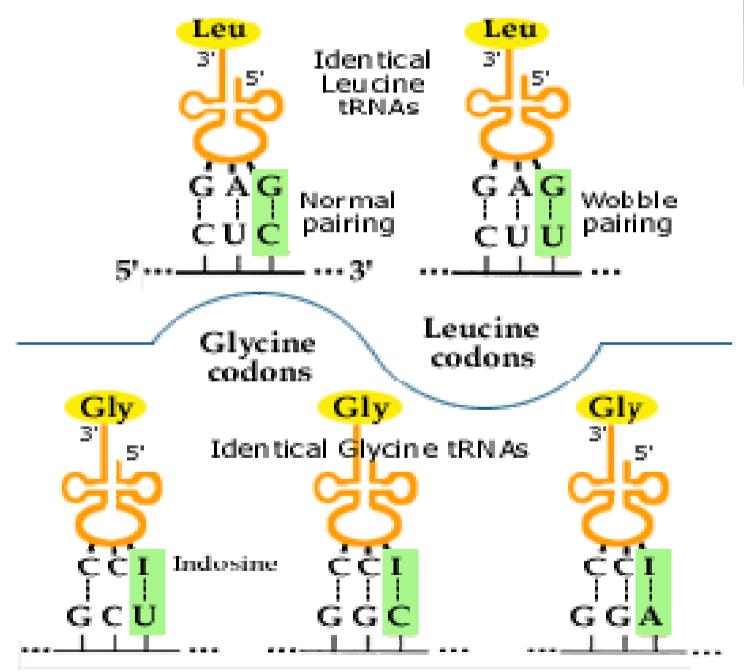
### 9. Wobbling Phenomenon:

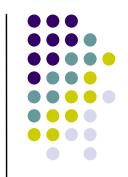
The reduced stringency between the third base of the codon and the complementary nucleotide in the anticodon is called wobble.

It reduce effect of mutation.

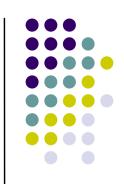
# **Wobbling Phenomenon**



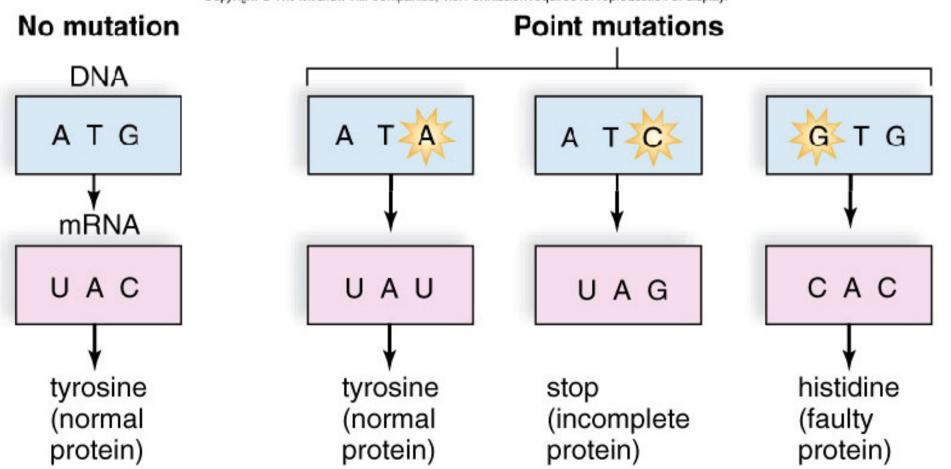


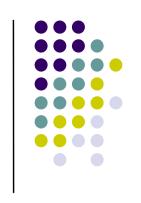


# Mutation



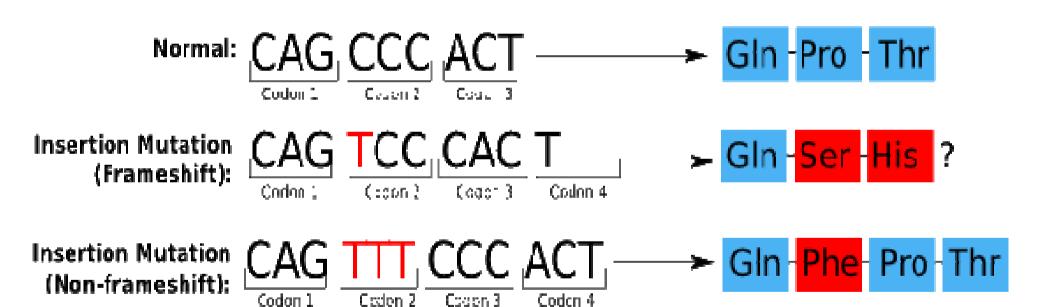
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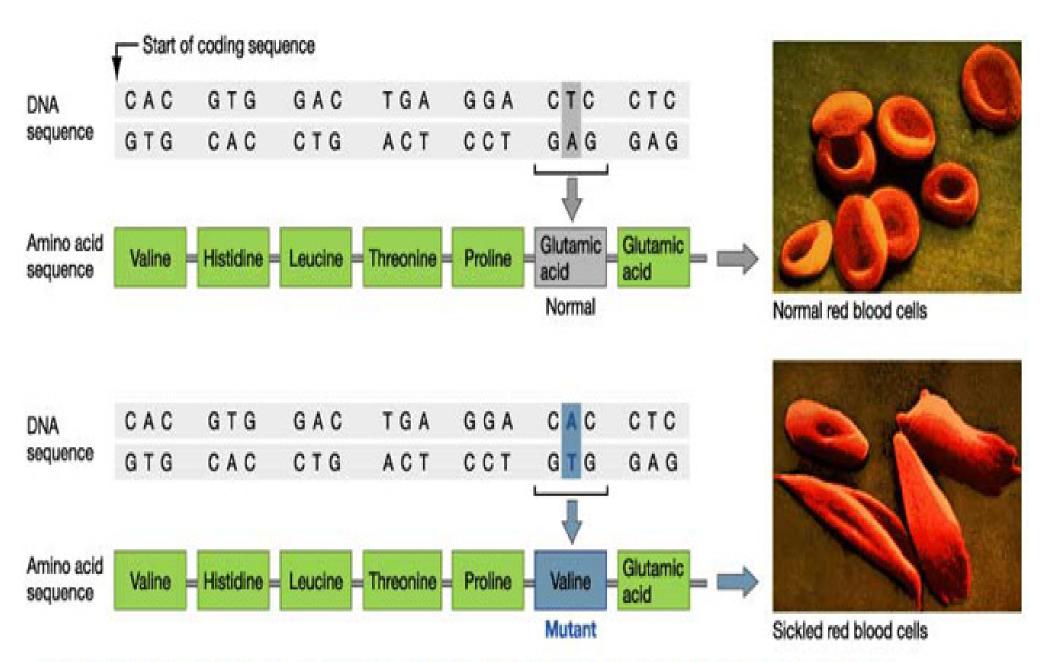




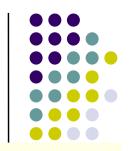
### **DNA Sequence**

### Amino Acid Sequence





The change in amino acid sequence causes hemoglobin molecules to crystallize when oxygen levels in the blood are low. As a result, red blood cells sickle and get stuck in small blood vessels.



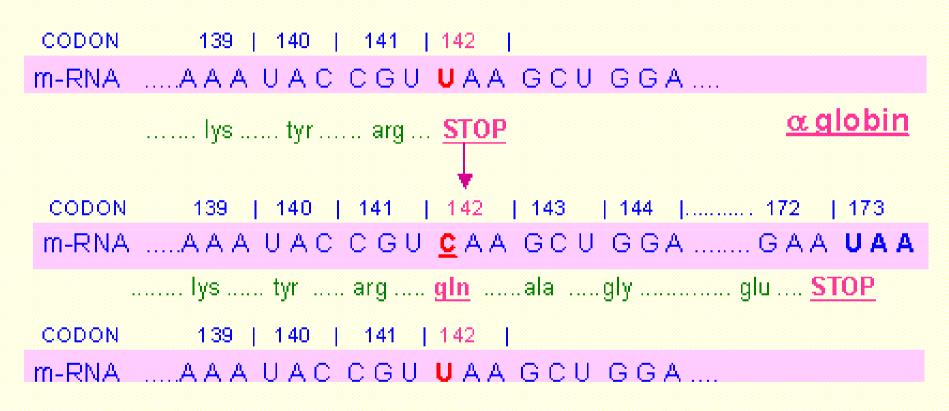
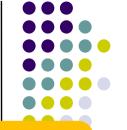


Fig 5.3. Point mutation of U to C alters the stop codon UAA at position 142, resulting in  $\alpha$ -globin chain elongation and formation of a variant haemoglobin. Hb Constant Spring.

### **Mutation**



**Point Mutation** 

**Effect of Mutation** 

**Substitution Mutation** 

**Deletion Mutation** 

**Insertion Mutation** 

# **Transition Mutation**

Purine to Purine

# Transversion Mutation

Purine to Pyrimidine

#### **Gene Deletion**

Thalassemia

### **Codon Deletion**

Cystic fibrosis

#### **Base deletion**

Frame Shift Mutation

### **Gene Insertion**

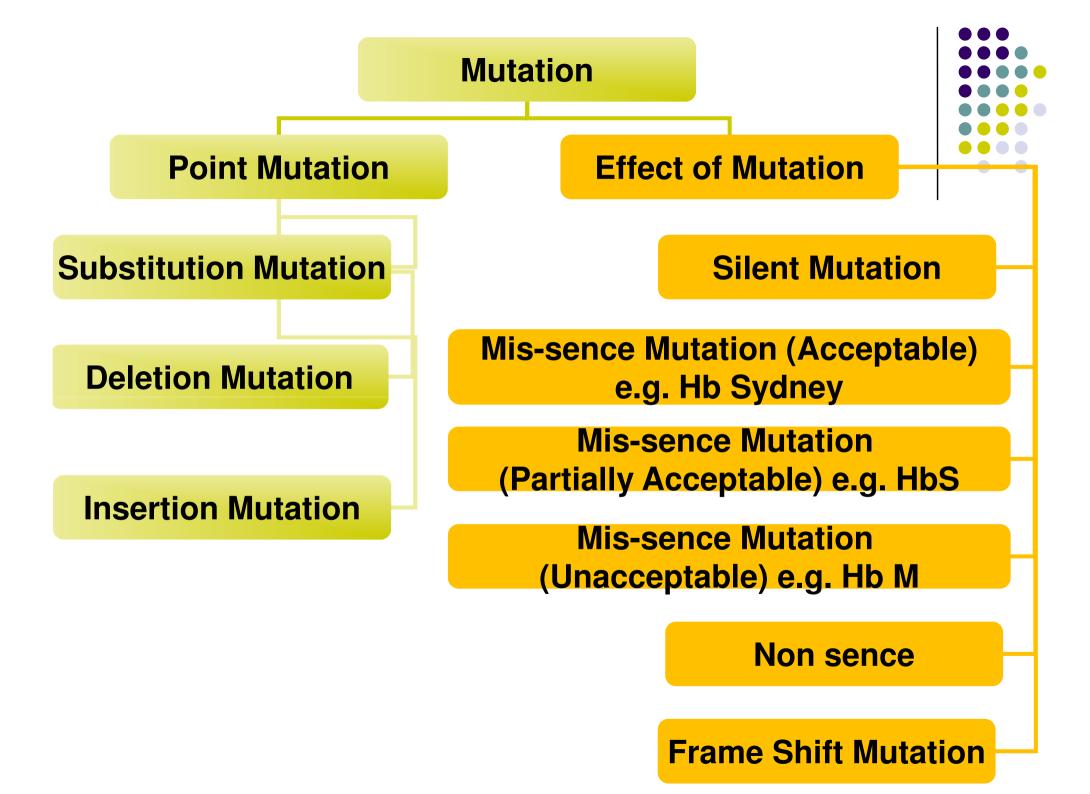
Duchenne Muscular Dystrophy

#### **Codon Insertion**

Huntington's chorea

### **Base deletion**

Frame Shift Mutation



# **Substitution Mutation**

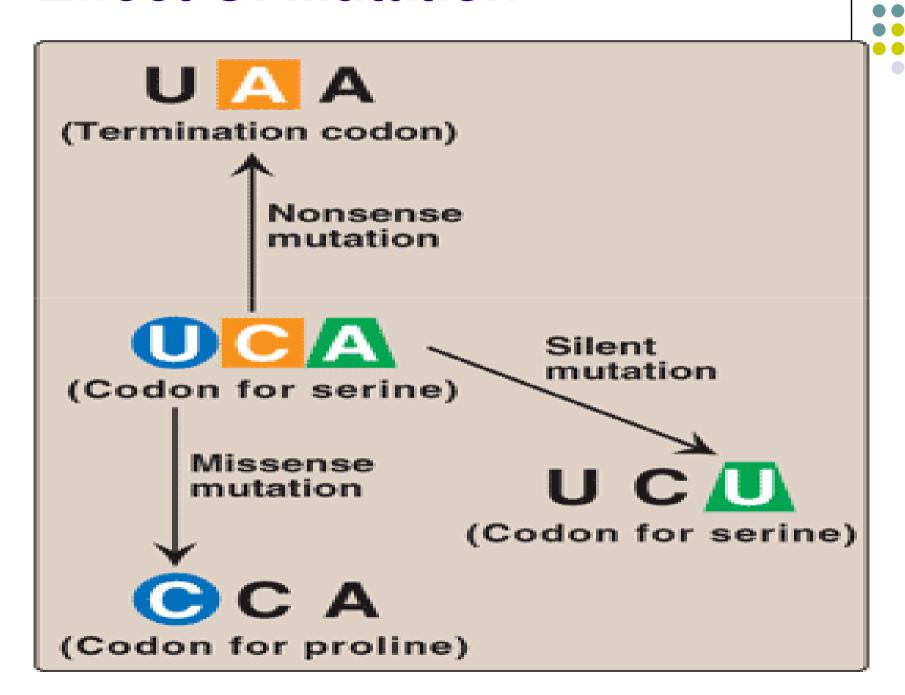


- Transition
  - Purine replace by Purine or
  - Pyrimidine replace by Pyrimidine

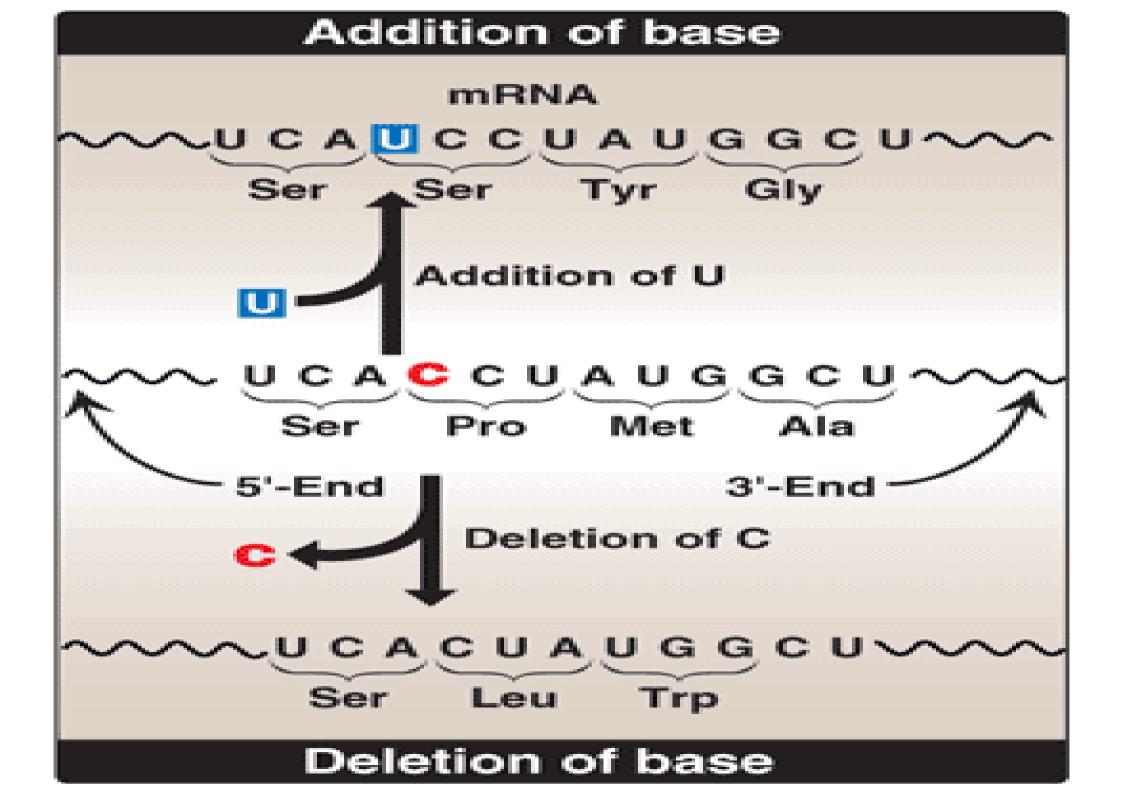
### Transversion

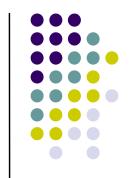
- Purine replace by Pyrimidine Or
- Pyrimidine replace by Purine
- E.g. Sickle cell anaemia
- GAG (glutamic acid)= GUG (valine)

# **Effect Of Mutation**



## **Huntington disease Huntington disease** / \ $(CAG)_{11-34}$ Tandem repeats of CAG triplets coding for glutamine mRNA is translated into huntinatin protein with abnormal glutamine repeats. Aggregated proteins and peptides Other triplet expansion diseases Fragile-X syndrome $(CGG)_{7-50}$ Myotonic dystrophy (CUG)5-35





# Translation (Protein Synthesis)

# **Translation**

- Steps:
  - Intiation
  - Elongation
  - Termination
  - Post Translation

# **Translation Process**



### Requires

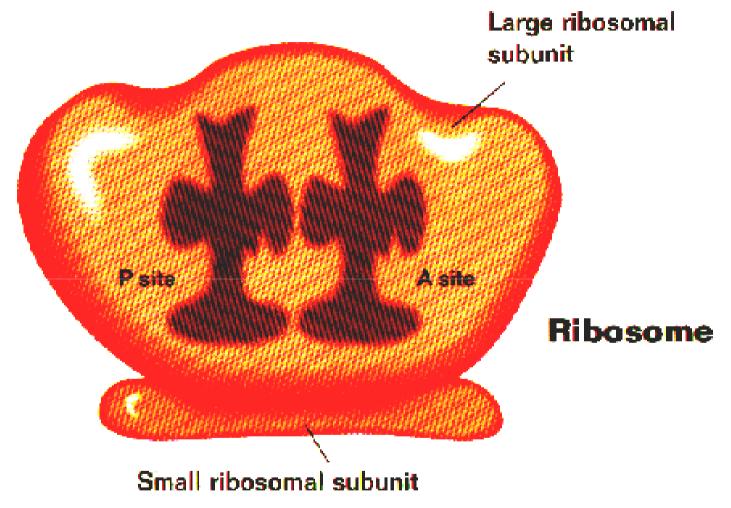
- Ribosomes (50s + 30s)
- r-RNA
- t-RNA
- m-RNA
- Amino acid

### Ribosome

- Made of protein and r-RNA (Nucleo-protein)
- Has internal sites for 2 t-RNA molecules.
- Two subunit
  - Prokaryotic 50S + 30S subunits = form a 70S.
  - Eukaryotic 60S + 40S subunits = form an 80S.

# Ribosome





## **Translation**

### Initiation

- Ribosomal subunits assemble on mRNA
- r-RNA aids in binding of mRNA



# Elongation

- t-RNAs with appropriate anticodon loops bind to complex
- have amino acid attached (done by other enzymes)
- Amino acids transfer form t-RNA 2 to t-RNA 1
- Process repeats

### Termination

- t-RNA with stop codon binds into ribosome
- No amino acid attached to t-RNA
- Complex falls apart

### A – P site on ribosome

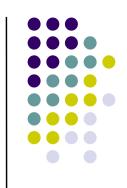
- Ribosome has two binding sites for t-RNA
  - P & A sites —
- Together, they cover two neighboring codons.



- codon is occupied by Peptidyl-tRNA.
- This tRNA carries the chain of amino acids that has already been synthesized.

### A site binds

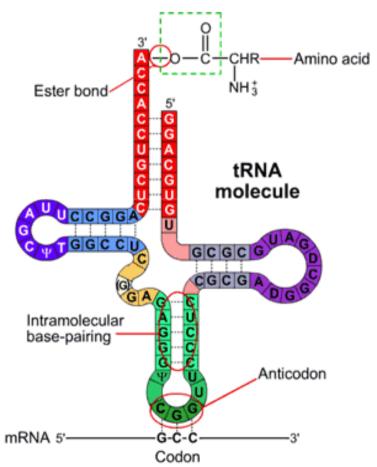
- incoming Aminoacyl-tRNA as directed by the codon.
- This codon specifies the next amino acid to be added to the growing peptide chain.

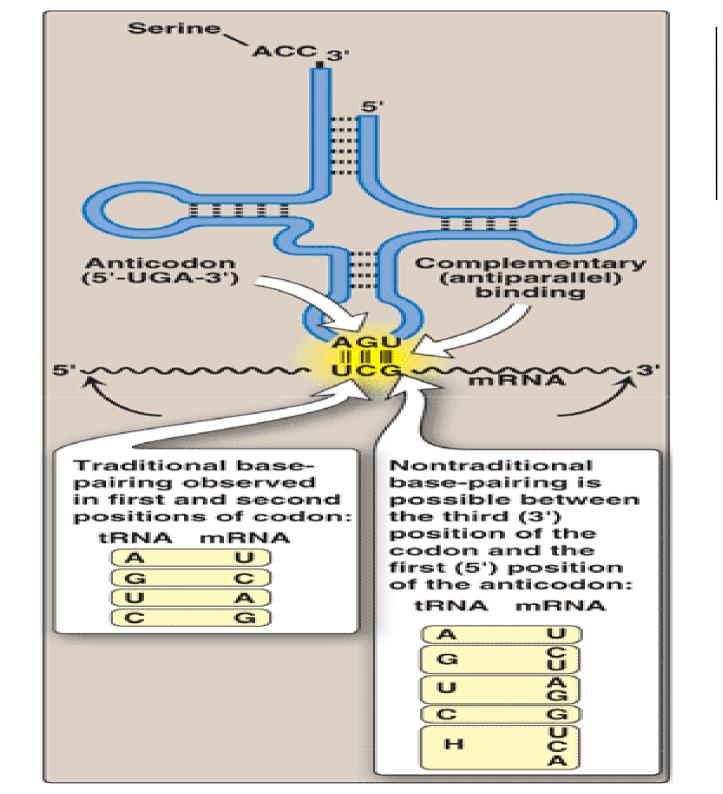


# **Transfer RNA**



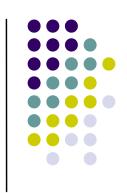






# **Aminoacyl-tRNA synthetases**

 Required for attachment of amino acids to their corresponding t-RNA.



- Carboxyl group of an amino acid to the 3'-end of t-RNA.
- Require ATP.
- Extreme specificity
- High fidelity of translation of the genetic message.
- Enzyme also have a "proofreading" or "editing" activity that can remove mischarged amino acids from the enzyme or the t-RNA molecule.

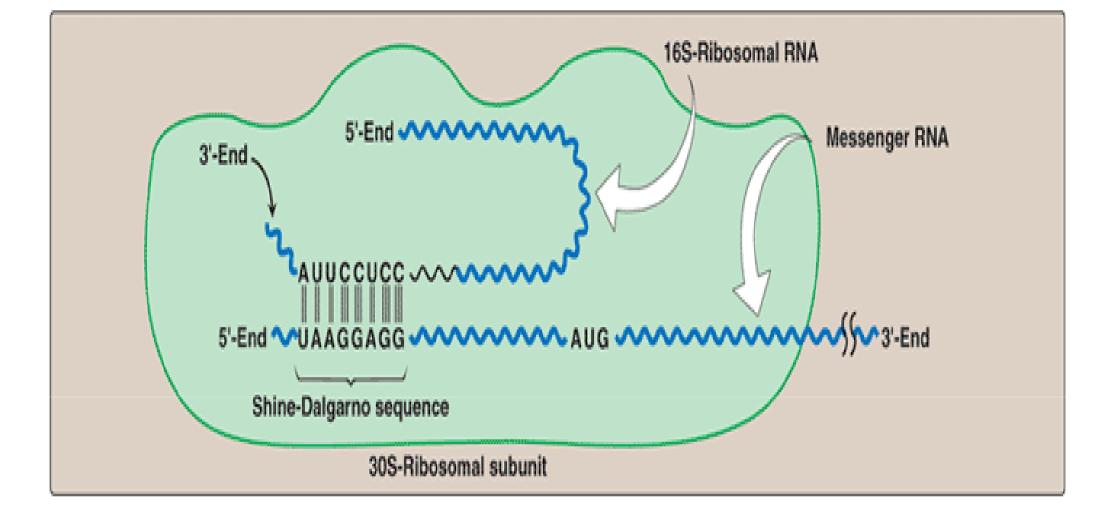
## **Initiation**



- Involves the assembly of the components of the translation system before peptide bond formation occurs.
- Components include
  - Two ribosomal subunits.
  - m-RNA to be translated.
  - Aminoacyl t-RNA specified by the first codon in the message
  - GTP
  - Initiation factors
    - In Prokaryotes, three initiation factors (IF-1, IF-2, IF-3)
    - In eukaryotes, there are over ten (designated eIF).

### Shine - Dalgarno (SD) sequence

- Purine rich sequence (e.g. 5'-UAAGGAGG-3')
- 6 10 bases upstream of the initiating AUG codon
- Near m-RNA 5'-end.
- 16S r-RNA of the 30S ribosomal subunit has complementary base pair at 3'-end.
- Facilitating the binding of the 30S ribosomal subunit on the mRNA
- In eukaryotes the 40S ribosomal subunit binds to the cap structure at the 5'-end of the mRNA by eIF-4 and moves down the mRNA until it encounters the initiator AUG. This "scanning" process requires ATP.

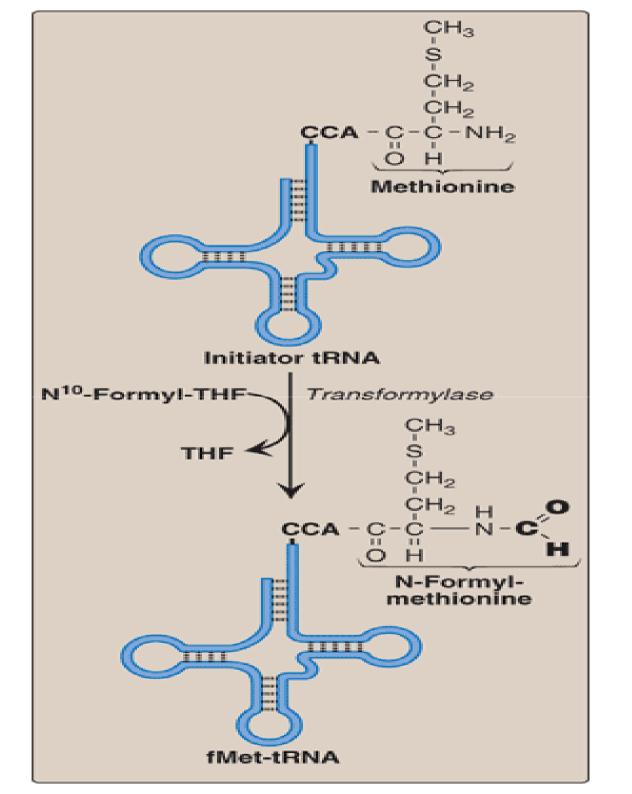


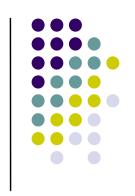
Complementary binding between prokaryotic mRNA Shine-Dalgarno sequence and 16S rRNA.

### **Initiation codon**

- AUG is recognized by initiator t-RNA.
- Recognition is facilitated by IF-2 (bound to GTP).

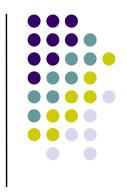
- Initiator t-RNA enters the ribosomal P site
- GTP is hydrolyzed to GDP.
- Initiator t-RNA carries N-formylated methionine.
- N10-formyl tetrahydrofolate as the carbon donor.
- In eukaryotes, the initiator tRNA carries a methionine that is not formylated.

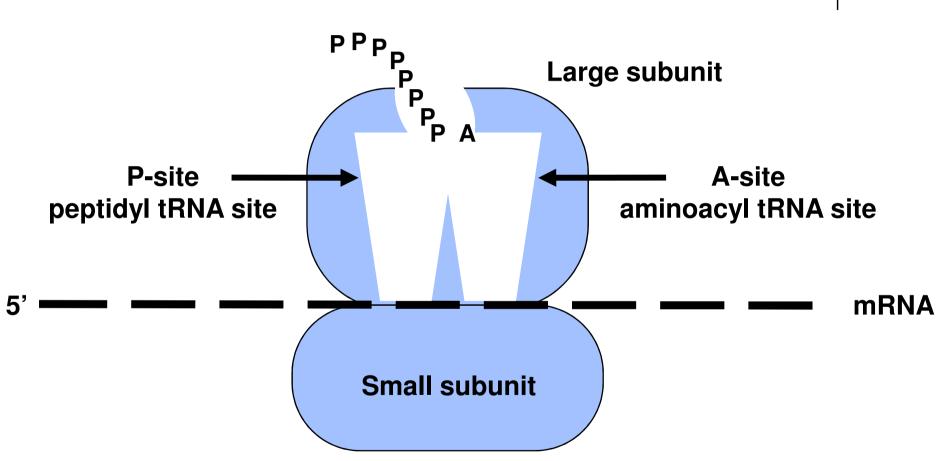




Generation of the initiator
N-formylmethionyl-tRNA
(fMet-tRNA)

#### Ribosome structure





Ribosome with bound tRNAs and mRNA

#### Initiation of protein synthesis: mRNA binding



M

Initiator tRNA bound to the small ribosomal subunit with the eukaryotic initiation factor-2 (eIF2)

elF2

40S subunit

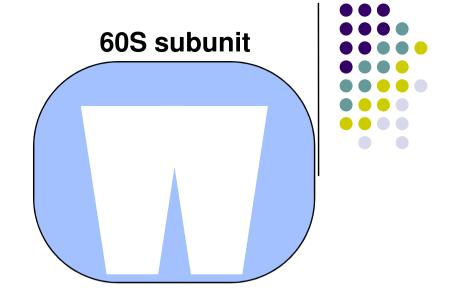
The small subunit finds the 5' cap and scans down the mRNA to the first AUG codon

5' cap —— —— —— AUG — —— mRNA

- the initiation codon is recognized
- eIF2 dissociates from the complex
- the large ribosomal subunit binds

elF2

M



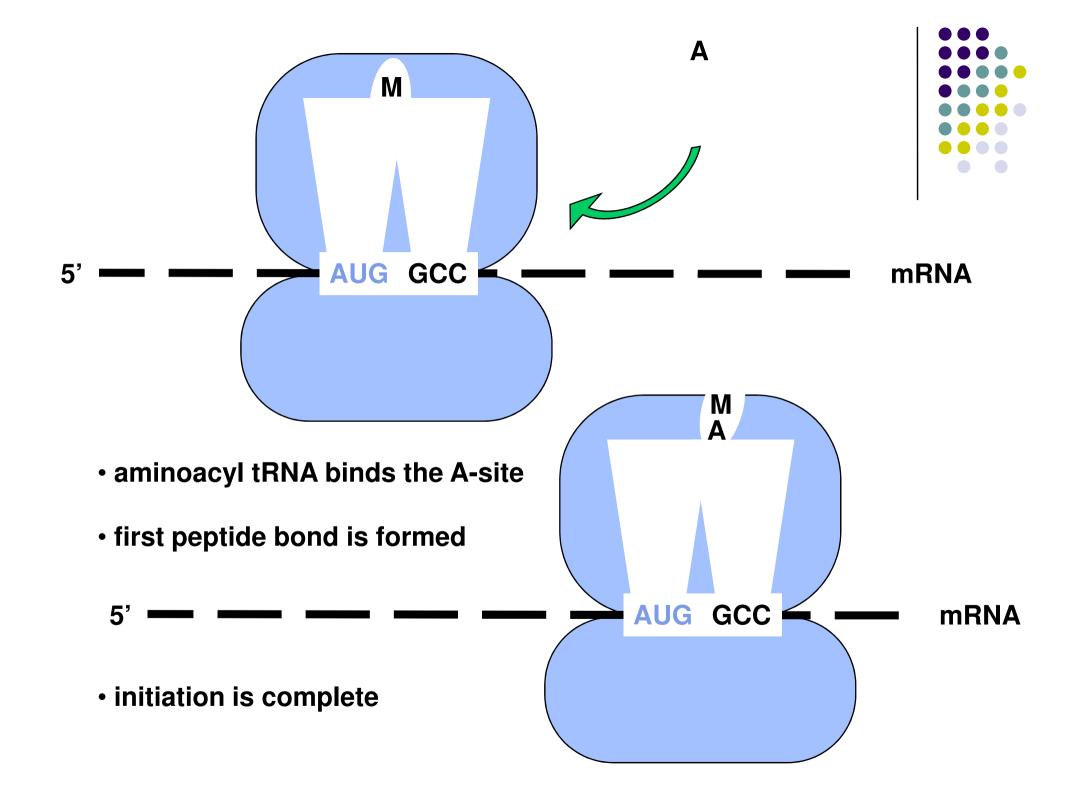
5' AUG MRNA
40S subunit

# Elongation

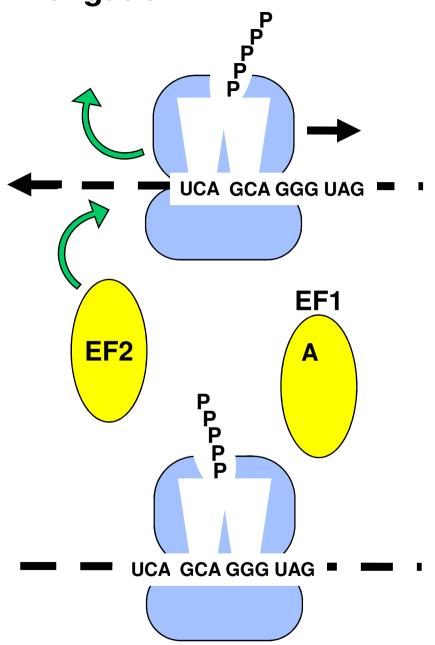
 Addition of amino acids to the carboxyl end of the growing chain.



- Ribosome moves from the 5'-end to the 3'-end of the mRNA that is being translated.
- New aminoacyl-tRNA for next triplet codon appears in ribosomal A site is facilitated in by EF-Tu, EF-Ts & EF-G and requires GTP hydrolysis.
- After the peptide bond has been formed, the ribosome moves to next triplet codon toward the 3'-end of the mRNA. This process is known as <u>Translocation</u>
- This causes movement of the uncharged tRNA into the ribosomal E site and movement of the peptidyl-tRNA into the P site.



#### **Elongation**



 the uncharged tRNA dissociate from the P-site



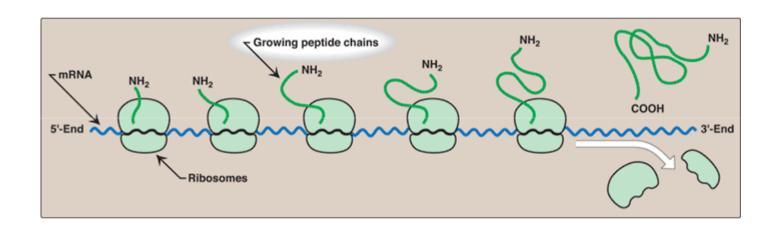
translocation =

the ribosome shifts one codon along mRNA, moving peptidyl tRNA from the A-site to the P-site.
requires EF2

 next aminoacyl tRNA then binds within the A-site; tRNA binding requires EF1

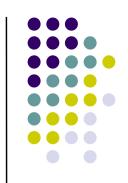
- energy for elongation is provided by the hydrolysis of two GTPs:
  - one for translocation
  - one for aminoacyl tRNA binding





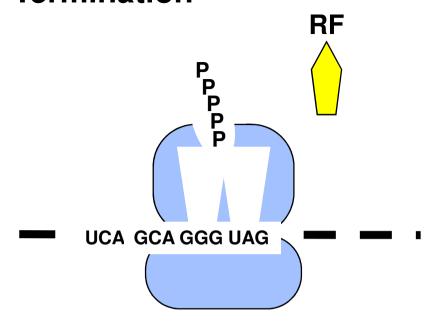
### **Termination**

 Termination occurs when one of the three termination codons moves into the A site.

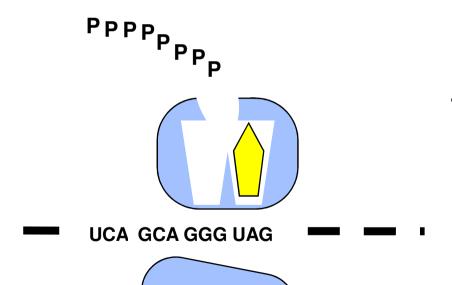


- Termination codons are recognized in by
  - RF-1 = UAA and UAG
  - RF-2 = UAA and UGA
- RF induces peptidyltransferase to hydrolyze the bond linking the peptide to the tRNA at the A site, causing the nascent protein to be released from the ribosome.
- RF-3 (bound to GTP) cause release of RF-1 or RF-2 as GTP is hydrolyzed.

#### **Termination**



 when translation reaches the stop codon, RF binds within the A-site, recognizing the stop codon



RF catalyzes the hydrolysis
 of the completed polypeptide from
 the peptidyl tRNA, and the entire
 complex dissociates

# Post-translation modification Trimming

- Many proteins are initially made as initially made as large, precursor molecules that are not functionally active.
- Precursor proteins are cleaved in the endoplasmic reticulum or the Golgi apparatus, others are cleaved in developing secretory vesicles.
- Zymogens are inactive secreted enzymes (including the proteases required for digestion).
- They become activated through cleavage when they reach their proper sites of action.
- E.g. Pancreatic zymogen, trypsinogen, becomes activated to in the small intestine.

## Post-translation modification

#### **Covalent Alterations**

### 1. Phosphorylation:

- On hydroxyl groups of serine, threonine, tyrosihe.
- increase or decrease the functional activity of the protein.

## 2. Glycosylation:

- Proteins of a plasma membrane.
- Carbohydrate attached to serine or threonine hydroxyl groups (O-linked) or the amide nitrogen of asparagine (N-linked).

### 3. Hydroxylation:

Proline and lysine residues = collagen

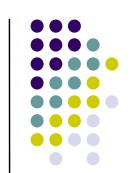


### Post-translation modification

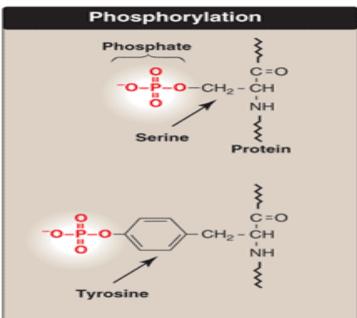
#### **Covalent Alterations**

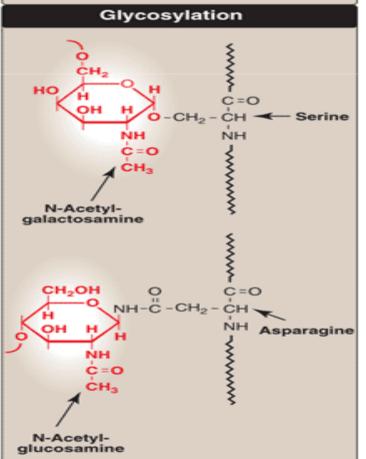
### 1.Other covalent modification:

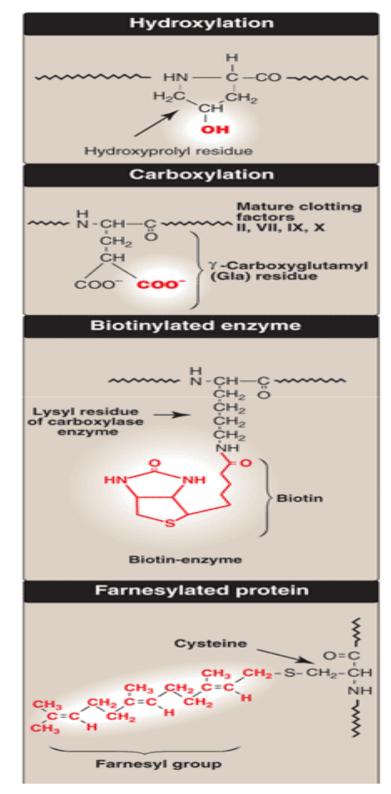
 Vitamin K-dependent carboxylation of Glutamate residues of clotting factors.



- Biotin = ε-amino groups of lysine residues of biotin-dependent enzymes
  - = carboxylation reactions.
  - e.g. pyruvate carboxylation
- Attachment of lipids, such as farnesyl groups, can help anchor proteins in membranes.
- Acetylated









## Post-translation modification

#### **Protein degradation**

- Defective Protein, for rapid turnover are often marked for destruction by ubiquitination
- The attachment of a small, highly conserved protein, called ubiquitin.
- Proteins marked in this way are rapidly degraded by a cellular component known as the "proteasome".

# **Inhibitor of Translation**



<u>Inhibitor ( Antibiotic)</u>	
Erythromycin	50 s ribosomal subunit
Clindamycin	50 s ribosomal subunit
Tetracycline	30 s ríbosomal subunít
Puramycin	Amino-acyl t-RNA
Chloramphenicol	Peptidyl Transferase
Diphtheria Toxin	eEF-2

